WE CLAIM:

- In a method of detecting a plurality of electrophoretically separated classes of DNA
 fragments, an improvement comprising labelling DNA fragments of at least one class with a 4,7-dichlorofluorescein dye.
 - 2. The method of claim 1 wherein said 4,7-dichlorofluorescein dye is defined by the formula:

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wherein:

A' is hydrogen, fluoro, chloro, a linking functionality, or a group that may be converted to a linking functionality;

B' is fluoro, chloro, or an acidic anionic group;

X' is hydrogen, fluoro, or chloro;

Z₁ is hydrogen or, when taken with Z₂, benzo;

Z2, when taken alone, is hydrogen, halo, lower alkyl, lower alkyloxy, a linking functionality, or a group that may be converted to a linking functionality, or when taken with Z1, benzo;

Z3 and Z4 are separately hydrogen, halo, lower alkyl, lower alkyloxy, a linking

functionality, or a group that may be converted to a linking functionality;

Z5, when taken alone, is hydrogen, halo, lower alkyl, lower alkyloxy, a linking functionality, or a group that may be converted to a linking functionality, or when taken with Z6, benzo;

Z6 is hydrogen or, when taken with Z5, benzo; and wherein at least one of A, Z2, Z3, Z4, and Z5 is a linking functionality or a group that may be converted to a linking functionality.

3. The method of claim 2 wherein:

A' is carboxyl, sulfonyl, isothlocyanate, succinimidyl carboxylate, phosphoramidite, or amino:

B' is carboxyl or sulfonyl;

X' is hydrogen or chloro;

Z2, when taken alone, is hydrogen, methyl, ethyl, methoxy, ethoxy, or chloro;

Z3, and Z4 are separately hydrogen, methyl, ethyl, methoxy, ethoxy, chloro, carboxyl,

sulfonyl, isothiocyanate, succinimidyl carboxylate, phosphoramidite, or methylamino;

Z5, when taken alone, is hydrogen, methyl, ethyl, methoxy, ethoxy, or chloro; and wherein only one of A', Z3, and Z4 is carboxyl, sulfonyl, methylamino, isothiocyanate, succinimidyl carboxylate, phosphoramidite, or amino.

- 20 4. The method of claim 3 wherein Z3, and Z4 are separately hydrogen, methyl, ethyl, methoxy, ethoxy, or chloro.
 - 5. The method of claim 4 wherein Z2, when taken alone, is hydrogen, methoxy, ethoxy, or chloro;

Z3, and Z4 are separately hydrogen, methoxy, ethoxy, chloro; and Z5, when taken alone, is hydrogen, methoxy, ethoxy, or chloro.

6. The method of claim 5 wherein B' is carboxy and A' is carboxy, succinimidyl carboxylate, or phosphoramidite.

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7. A compound having the formula:

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wherein:

A' is hydrogen, fluoro, chloro, a linking functionality, or a group that may be converted to a linking functionality;

B' is fluoro, chioro, or an acidic anionic group;

X' is hydrogen, fluoro, or chioro;

Z3 and Z4 are separately hydrogen, halo, a linking functionality, or a group that may be converted to a linking functionality; and

wherein at least one of A', Z3, and Z4 is a linking functionality or a group that may be converted to a linking functionality.

- 8. The compound of claim 7 wherein A' is carboxyl, sulfonyl, isothiocyanate, succinimidyl carboxylate, phosphoramidite, or amino; B' is carboxyl or sulfonyl; X' is hydrogen; Z3 and Z4 are separately hydrogen, halo, carboxyl, sulfonyl, or methylamino.
- 9. The compound of claim 8 wherein only one of A', Z3, and Z4 is carboxyl, sulfonyl, methylamino, or amino.
- 35 10. The compound of claim 9 wherein A' and B' are carboxyl, Z3 is hydrogen or chloro, and

Z4 is hydrogen or chloro.

- 11. A kit for detecting a plurality of electrophoretically separated classes of DNA fragments comprising an oligonucleotide labelled with a 4,7-dichlorofluorescein dye.
- 12. The kit of claim 11 further comprising:

an enzyme selected from the group consisting of nucleic acid polymerase and nucleic acid ligase; and

a reaction buffer.

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- 13. The kit of claim 12 wherein said enzyme is a nucleic acid polymerase and wherein said kit further includes a nucleoside triphosphate mix.
- 14. The kit of claim 12 wherein said enzyme is a nucleic acid ligase.

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- 15. A kit for sequenceing DNA comprising:
 - an oligonucleotide labelled with a 4,7-dichlorofluorescein dye:
 - a nucleic acid polymerase;
 - a reaction buffer; and
- 20 a nucleoside triphosphate mix.
 - 16. The kit of claim 15 wherein said 4,7-dichlorofluorescein dye is defined by the formula:

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wherein:

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A' is hydrogen, fluoro, chloro, or a group that may be converted to a linking functionality;

B' is fluoro, chloro, or an acidic anionic group:

X is hydrogen, fluoro, or chloro;

Z₁ is hydrogen or, when taken with Z₂, benzo:

Z2, when taken alone, is hydrogen, halo, lower alkyl, lower alkyloxy, or a group that may be converted to a linking functionality, or when taken with Z1, benzo;

Z3 and Z4 are separately hydrogen, halo, lower alkyl, lower alkyloxy, or a group that may be converted to a linking functionality:

Z5, when taken alone, is hydrogen, halo, lower alkyl, lower alkyloxy, or a group that may be converted to a linking functionality, or when taken with Za, benzo;

Za is hydrogen or, when taken with Z5, benzo; and wherein at least one of A', Z2, Z3, Z4, and Z5 is a group that may be converted to a linking functionality.

15 17. The kit of claim 16 wherein:

A' and B' are carboxyl:

X' is hydrogen or chloro;

Z2, when taken alone, is hydrogen, methyl, ethyl, methoxy, ethoxy, or chloro;

Z3, and Z4 are separately hydrogen, methyl, ethyl, methoxy, ethoxy, chloro; and

Z5, when taken alone, is hydrogen, methyl, ethyl, methoxy, ethoxy, or chloro.

- 18. A kit for sequencing DNA comprising:
- a dye-terminator mix wherein at least one dye-terminator is labelled with a 4.7dichlorofluorescein dye:
 - a nucleic acid polymerase:
 - a nucleoside triphosphate mix; and
 - a reaction buffer.
- 19. The kit of claim 18 wherein said dye-terminator mix comprises dideoxynucleoside 30 triphosphates selected from the group consisting of dideoxyadenosine triphosphate, dideoxycytidine triphosphate, dideoxyguanosine triphosphate, and dideoxythymidine triphosphate wherein each of said dideoxynucleoside triphosphates is separately labelled with a dye selected from the group consisting of 5- and 6-carboxyfluorescein, 5- and 6-carboxy-4,7dichlorofluorescein, 2',7'-dimethoxy-5- and 6-carboxy-4,7-dichlorofluorescein, 2',7'-dimethoxy-
- 35 4',5'-dichloro-5- and 6-carboxyfluorescein, 2',7'-dimethoxy-4',5'-dichloro-5- and 6-carboxy-4,7-

dichlorofluorescein, 1',2',7',8'-dibenzo-5- and 6-carboxy-4,7-dichlorofluorescein, 1',2',7',8'-dibenzo-4',5'-dichloro-5- and 6-carboxy-4,7-dichlorofluorescein, 2',7'-dichlorofluorescein, and 2',4',5',7'-tetrachloro-5- and 6-carboxy-4,7-dichlorofluorescein.

- 5 20. The kit of claim 19 wherein said dideoxythymidine triphosphate is labelled with 6-carboxyfluorescein, said dideoxycytidine triphosphate is labelled with 2',4',5',7'-tetrachloro-5-carboxyfluorescein, said dideoxyadenosine triphosphate is labelled with 2',4',5',7'-tetrachloro-4,7-dichloro-5-carboxyfluorescein, and said dideoxyguanosine triphosphate is labelled with 1',2',7',8'-dibenzo-4,7-dichloro-5-carboxyfluorescein.
 - 21. The kit of claim 20 wherein said nucleic acid polymerase is Sequenase TM.

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